

department of the four hospital sites with febrile illness, and will be interviewed to gather information on symptoms, possible exposures, and medical history, in addition to having diagnostic samples collected to test for leptospirosis plus or minus melioidosis (depending on presenting symptoms).

Participants will then also be interviewed approximately two weeks after enrollment to determine illness progression and outcome. Patients testing positive for leptospirosis, if willing, may have animals sampled from their home or work environments, if present, to help determine the animal

reservoirs related to human leptospirosis illness in Puerto Rico.

The estimated annualized burden hours requested are 1,675. There is no cost to respondents other than their time.

#### ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Patient .....	PIFA (screening) .....	7,000	1	5/60	583
Patient .....	PIFA (full form: Sections 1–4, 11) ...	3,000	1	10/60	500
Patient .....	Consent Form .....	3,000	1	6/60	300
Patient .....	PIFF .....	1,000	1	10/60	167
Patient .....	Animal Household Survey .....	250	1	30/60	125
Total .....	.....	.....	.....	.....	1,675

**Jeffrey M. Zirger,**

*Lead, Information Collection Review Office,  
Office of Scientific Integrity, Office of Science,  
Centers for Disease Control and Prevention.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92–463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)–CE15–002: The CDC National Centers of Excellence in Youth Violence Prevention: Building the Evidence for Community- and Policy-Level Prevention.

*Date:* June 17, 2020.

*Time:* 8:30 a.m., EDT.

*Place:* Videoconference.

*Agenda:* To review and evaluate grant applications.

*For Further Information Contact:*  
Mikel Walters, Ph.D., Scientific Review Official, NCIPC, CDC, 4770 Buford Highway NE, Mailstop F–63, Atlanta, Georgia 30341, Telephone (404)639–0913, [MWalters@cdc.gov](mailto:MWalters@cdc.gov).

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

**Kalwant Smagh,**

*Director, Strategic Business Initiatives Unit,  
Office of the Chief Operating Officer, Centers  
for Disease Control and Prevention.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Notice of Change in Reporting Requirements for The Federal Cigarette Labeling and Advertising Act (FCLAA) and Comprehensive Smokeless Tobacco Health Education Act (CSTHEA)

**AGENCY:** Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** The Centers for Disease Control and Prevention (CDC), located within the Department of Health and Human Services (HHS), announces that it is extending the March 31, 2020 deadline for submissions required under the Federal Cigarette Labeling and Advertising Act (FCLAA) and the Comprehensive Smokeless Tobacco Health Education Act (CSTHEA) for cigarette and smokeless tobacco products, respectively. CDC also announces that it is extending the deadline for ingredient reports for new products that are due at the time of first importation. Previous notices announced that ingredient reports are due annually on March 31, and/or upon initial importation of cigarettes and/or smokeless tobacco products. Due to the current public health response to COVID–19, CDC is not able to accept any ingredient submissions and will not be issuing Certificates of Compliance at this time. CDC is communicating this information to state government entities and will re-evaluate this approach as necessary.